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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/830,147	04/22/2004	Martin Friedrich Stefanic	1/1464US	7845
28501 7590 04/03/2007 MICHAEL P. MORRIS BOEHRINGER INGELHEIM CORPORATION 900 RIDGEBURY ROAD P. O. BOX 368 RIDGEFIELD, CT 06877-0368			EXAMINER	
			ANDERSON, JAMES D	
			ART UNIT	PAPER NUMBER
			1614	
SHORTENED STATUTORY P	PERIOD OF RESPONSE	MAIL DATE	DELIVERY MODE	
31 DAYS 04/03/2007			PAPED	

# Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

Paper No(s)/Mail Date

Information Disclosure Statement(s) (PTO/SB/08)

Notice of Informal Patent Application

Other:

Application/Control Number: 10/830,147

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## **DETAILED ACTION**

#### Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. § 121:

- I. Claims 1-18, drawn to pharmaceutical combinations, classified in class 514, subclass 418. <u>Please note additional Election of Species requirement outlined</u> below.
- II. Claims 19-21, drawn to pharmaceutical combination preparation kits, classified in class 514, subclass 418. <u>Please note additional Election of Species requirement outlined below.</u>
- III. Claims 22-30, drawn to methods of treatment, classified in class 514, subclass418. Please note additional Election of Species requirement outlined below.

The inventions are distinct, each from the other because of the following reasons:

Inventions I/II and III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product. See MPEP § 806.05(h). In the instant case the methods of Group III recite the treatment of diseases involving cell proliferation, migration or apoptosis of myeloma cells or angiogenesis. However, the compositions recited in the claims of Groups I and II could be used in *in vitro* screening assays of any number of receptors (e.g., VEGF, PDGFR, HER2, etc.). As such, the products of Groups I and II could be used in a materially different process of using those products than the *in vivo* methods recited in the claims of Group III.

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Inventions I and II are directed to related pharmaceutical combinations. The related inventions are distinct if the (1) the inventions as claimed are either not capable of use together or can have a materially different design, mode of operation, function, or effect; (2) the inventions do not overlap in scope, i.e., are mutually exclusive; and (3) the inventions as claimed are not obvious variants. See MPEP § 806.05(j). In the instant case, the inventions as claimed have materially different designs and do not overlap in scope. For example, the claims of Group II require that the antagonist is comprised in a first compartment and the further chemotherapeutic agent is comprised within a second compartment. The combinations recited in the claims of Group I do not require this limitation. Furthermore, the inventions as claimed do not encompass overlapping subject matter and there is nothing of record to show them to be obvious variants.

These inventions are independent or distinct for the reasons given above and there would be a serious burden on the examiner if restriction is not required. Because the inventions have acquired a separate status in the art due to their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

### Election of Species Requirement Upon Election of Group I, II or III

Claims 1-30 are generic to the following disclosed patentably distinct species: A) the multitude of structurally unrelated antagonists of at least one receptor selected from VEGFR, PDGFR, FGFR, etc. and B) the multitude of structurally unrelated chemotherapeutic agents. The species are independent or distinct because the antagonists of at least one receptor selected from VEGFR, PDGFR, FGFR, etc. are structurally unrelated and would require different searches.

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Similarly, the term "chemotherapeutic" is generic and encompasses a multitude of agents with different classifications, mechanisms of action, and structures. Applicant is required under 35 U.S.C. § 121 to elect a single disclosed species, even though this requirement is traversed. Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

To be fully responsive to this Election of Species Requirement, Applicants must identify and elect a <u>single</u> antagonist of at least one receptor selected from VEGFR, PDGFR, FGFR, etc. and a <u>single</u> chemotherapeutic.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which depend from or otherwise require all the limitations of an allowable generic claim as provided by 37 CFR § 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

#### Joint Inventors

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR § 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR § 1.48(b) and by the fee required under 37 CFR § 1.17(i).

### Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to James D. Anderson whose telephone number is 571-272-9038. The examiner can normally be reached on MON-FRI 9:00 am - 5:00 pm EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ardin Marschel can be reached on 571-272-0718. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

James D. Anderson, Ph.D.

Patent Examiner

AU 1614

March 30, 2007

PHYLLIS SPIVACK
PRIMARY EXAMINER